

REC'D 05 DEC 2005

## PATENT COOPERATION TREATY

WIPO

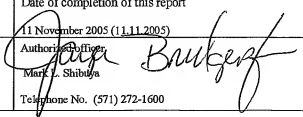
PCT

## PCT

## INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 10589-33-228	<b>FOR FURTHER ACTION</b>		See Form PCT/IPEA/416
International application No. PCT/US04/09574	International filing date ( <i>day/month/year</i> ) 26 March 2004 (26.03.2004)	Priority date ( <i>day/month/year</i> ) 27 March 2003 (27.03.2003)	
International Patent Classification (IPC) or national classification and IPC IPC(7): A01N 61/00; C12Q 1/00; G01N 33/566, 573 AND 574 AND US CL: 435/4, 6, 7.2, 7.21, 41, 69.2, 91.3, 183; 514/1, 2			
Applicant PTC THERAPEUTICS, INC.			
<p>1. This report is the international preliminary examination report, established by this International Preliminary Examining Authority under Article 35 and transmitted to the applicant according to Article 36.</p> <p>2. This REPORT consists of a total of <u>6</u> sheets, including this cover sheet.</p> <p>3. This report is also accompanied by ANNEXES, comprising:</p> <p>a. <input type="checkbox"/> (<i>sent to the applicant and to the International Bureau</i>) a total of ___ sheets, as follows:</p> <p><input type="checkbox"/> sheets of the description, claims and/or drawings which have been amended and are the basis of this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions).</p> <p><input type="checkbox"/> sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box.</p> <p>b. <input type="checkbox"/> (<i>sent to the International Bureau only</i>) a total of (indicate type and number of electronic carrier(s)) _____, containing a sequence listing and/or tables related thereto, in electronic form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the Administrative Instructions).</p> <p>4. This report contains indications relating to the following items:</p> <p><input checked="" type="checkbox"/> Box No. I Basis of the report</p> <p><input type="checkbox"/> Box No. II Priority</p> <p><input type="checkbox"/> Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability</p> <p><input checked="" type="checkbox"/> Box No. IV Lack of unity of invention</p> <p><input checked="" type="checkbox"/> Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability, citations and explanations supporting such statement</p> <p><input type="checkbox"/> Box No. VI Certain documents cited</p> <p><input type="checkbox"/> Box No. VII Certain defects in the international application</p> <p><input type="checkbox"/> Box No. VIII Certain observations on the international application</p>			
Date of submission of the demand 26 October 2004 (26.10.2004)		Date of completion of this report 11 November 2005 (11.11.2005)	
Name and mailing address of the IPEA/US Mail Stop PCT, Attn: IPEA/US Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450 Facsimile No. (571) 273-3201		 Authorized Officer Mark L. Shibuya Telephone No. (571) 272-1600	

Form PCT/IPEA/409 (cover sheet)(April 2005)

**Box No. I Basis of the report**

## 1. With regard to the language, this report is based on:

- ☒ the international application in the language in which it was filed.
- ☐ a translation of the international application into English, which is the language of a translation furnished for the purposes of:
- ☐ international search (under Rules 12.3 and 23.1(b))
- ☐ publication of the international application (under Rule 12.4(a))
- ☐ international preliminary examination (under Rules 55.2(a) and/or 55.3(a))

2. With regard to the elements of the international application, this report is based on (*replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report*):

- ☒ the international application as originally filed/furnished
- ☒ the description:  
pages 1-102 as originally filed/furnished  
pages\* NONE received by this Authority on \_\_\_\_\_  
pages\* NONE received by this Authority on \_\_\_\_\_
- ☒ the claims:  
pages 103-111 as originally filed/furnished  
pages\* NONE as amended (together with any statement) under Article 19  
pages\* NONE received by this Authority on \_\_\_\_\_  
pages\* NONE received by this Authority on \_\_\_\_\_
- ☒ the drawings:  
pages 1/1 as originally filed/furnished  
pages\* NONE received by this Authority on \_\_\_\_\_  
pages\* NONE received by this Authority on \_\_\_\_\_

☐ a sequence listing and/or any related table(s) - see Supplemental Box Relating to Sequence Listing.

3. ☒ The amendments have resulted in the cancellation of:

- ☒ the description, pages NONE
- ☒ the claims, Nos. NONE
- ☒ the drawings, sheets/figs NONE
- ☒ the sequence listing (*specify*): NONE
- ☒ any table(s) related to the sequence listing (*specify*): NONE

4. ☐ This report has been established as if (some of) the amendments annexed to this report and listed below had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(e)).

- ☐ the description, pages \_\_\_\_\_
- ☐ the claims, Nos. \_\_\_\_\_
- ☐ the drawings, sheets/figs \_\_\_\_\_
- ☐ the sequence listing (*specify*): \_\_\_\_\_
- ☐ any table(s) related to the sequence listing (*specify*): \_\_\_\_\_

\* If item 4 applies, some or all of those sheets may be marked "superseded."

## INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/US04/09574

**Box No. IV Lack of unity of invention**

1. ☐ In response to the invitation to restrict or pay additional fees the applicant has, within the applicable time limit:
- ☐ restricted the claims.
  - ☐ paid additional fees.
  - ☐ paid additional fees under protest, and, where applicable, the protest fee
  - ☐ paid additional fees under protest but the applicable protest fee was not paid
  - ☐ neither restricted the claims nor paid additional fees
2. ☒ This Authority found that the requirement of unity of invention is not complied with and chose, according to Rule 68.1, not to invite the applicant to restrict or pay additional fees.
3. This Authority considers that the requirement of unity of invention in accordance with Rules 13.1, 13.2 and 13.3 is:
- ☐ complied with.
  - ☒ not complied with for the following reasons:

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Group I, claim(s) 1-28 and 33-39, drawn to methods for identifying a compound that modulates fungal tRNA splicing endonuclease activity.

Group II, claim(s) 29-32, 40 and 41, drawn to methods of preventing, treating, managing or ameliorating a fungal infection by administering an antiproliferative compound identified by the Group I method.

The inventions listed as Groups I and II do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: the methods of Groups I and II are distinctly different methods drawn to different method objectives. The antifungal compounds of Group II and derived from the Group I methods do not represent a "special" technical feature because antifungal compounds are known in the art. See e.g., WO 02/083953A1; WO 02/083837A1; and WO 01/25486A1.

4. Consequently, this report has been established in respect of the following parts of the international application:

- ☒ all parts
- ☐ the parts relating to claims Nos. \_\_\_\_\_

## INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.  
PCT/US04/09574**Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement**

## 1. Statement

Novelty (N)	Claims <u>1-28, 33-39</u>	YES
	Claims <u>29-32, 40, 41</u>	NO
Inventive Step (IS)	Claims <u>NONE</u>	YES
	Claims <u>1-41</u>	NO
Industrial Applicability (IA)	Claims <u>1-41</u>	YES
	Claims <u>NONE</u>	NO

2. Citations and Explanations (Rule 70.7)  
Please See Continuation Sheet

## Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

Continuation of:

## V. 2. Citations and Explanations:

Claims 29-32, 40 and 41 lack novelty under PCT Article 33(2) as being anticipated by US 5,726,195 A (HILL et al.).

Hill et al. discloses small molecule antifungal (e.g. anti-yeast) compounds for treating microbial infections when administered to a host, (e.g., human). These compounds inhibit tRNA enzymes (e.g. synthetases) and comprise structure within the scope of the presently claimed invention (e.g. see examples and patent claims). The ability to inhibit tRNA splicing endonuclease is inherently present due to the ability of these compounds to bind tRNA. In any event, the claim is not structure-limited and the PTO lacks the facilities for making comparisons between prior art compounds and the claimed prospective assay-derived compounds.

Claims 29-32, 40 and 41 lack novelty under PCT Article 33(2) as being anticipated by WO 01/25486 A1 (RANA).

Rana discloses assay-derived tRNA inhibiting (e.g., binding; see e.g. bottom of page 9-top of page 10; and claims, especially claims 1, 2, 28-30, 40-43) compounds within the scope of the presently claimed invention (e.g., claims 25-26) that are antifungal for use in treating fungal (e.g. yeast; see claims 47-48) infections (e.g., see page 10-11) when administered to humans. The ability to inhibit tRNA splicing endonuclease is inherently present due to the ability of these compounds to bind RNA (e.g. tRNA). In any event, the claim is not structure-limited and the PTO lacks the facilities for making comparisons between prior art compounds and the claimed prospective assay-derived compounds.

Claims 29-32, 40 and 41 lack novelty under PCT Article 33(2) as being anticipated by WO 02/083837 A1 (ALMSTEAD).

Almstead discloses assay-derived binding compounds (e.g. see pages 3-4; bottom of page 10-11) within the scope of the presently claimed invention (e.g. see pages 21-23; claim 5) that are antifungal for use in treating fungal (e.g., yeast) infections when administered to humans. The ability to inhibit tRNA splicing endonuclease is inherently present due to the ability of these compounds to bind RNA (e.g. tRNA). In any event, the claim is not structure-limited and the PTO lacks the facilities for making comparisons between prior art compounds and the claimed prospective assay-derived compounds.

Claims 29-32, 40 and 41 lack novelty under PCT Article 33(2) as being anticipated by WO 02/083953 A1 (RANDO et al.).

Rando et al. disclose assay-derived RNA binding (e.g., tRNA) compounds which effect RNA host cell factor complexes in vivo (e.g. RNA splicing; see page 10; bottom of page 12-page 13) which compounds are within the scope of the presently claimed invention (e.g. see claim 5) that are antifungal for use in treating fungal (e.g., yeast) infections when administered to humans. The ability to inhibit tRNA splicing endonuclease is inherently present due to the ability of these compounds to bind RNA (e.g. tRNA). In any event, the claim is not structure-limited and the PTO lacks the facilities for making comparisons between prior art compounds and the claimed

## Supplemental Box

prospective assay-derived compounds.

Claims 1-41 lack an inventive step under PCT Article 33(3) as being obvious over WO 01/25486 A1 (RANA), WO 02/083837 A1 (ALMSTEAD), and/or WO 02/083953 A1 (RANDO et al.) in view of WANG et al., *Nucleic Acids Research* Vol. 18, No. 22, HYDE-DERUYSCHER et al., *Chem. & Biol.* Vol. 7, No. 1, and LI et al., *Science* Vol. 280 (4/1999).

The presently claimed invention is directed to identifying antifungal compounds by screening (e.g., high throughput assays) compounds (e.g., library derived) for their ability to inhibit the endonucleolysis of fungal tRNA by inhibiting tRNA-tRNA splicing endonuclease binding, relative to a control.

Screening assays (e.g., high throughput assays) of single compounds or compound libraries for their ability to disrupt RNA (e.g., tRNA) interactions (e.g. including splicing) in order to identify antifungal drug candidates is taught by the RANA, ALMSTEAD and/or RANDO reference whose teaching discussed above is hereby incorporated by reference in its entirety.

The RANA, ALMSTEAD and/or RANDO reference methods differ from the presently claimed invention by failing to explicitly teach the application of its methods to tRNA splicing endonuclease assays that cleave tRNA and tRNA splicing endonuclease.

However, LI et al. teach that the tRNA splicing pathway is analogous in mammals and other organisms (e.g., fungi).

In this regard, WANG et al. teach an assay for endonucleolytic tRNA maturation, where inactivated micrococcal nuclease (reversible inhibitor) bound to radiolabeled pre-tRNA physically blocks the sites of endonuclease cleavage and prevents tRNA processing activities present in Fraction III of spinach chloroplasts, presumably by substrate occlusion or "masking", where formation of an inactive micrococcal nuclease enzyme substrate complex precludes utilization of the tRNA substrate by a second enzyme.

Additionally, the HYDE-DERUYSCHER et al. reference teaches that high throughput screening of "small molecule" compound libraries (e.g., phage) is ideal for screening "small molecule" enzyme inhibitors for a variety of different enzymes.

Accordingly, it would have been obvious to use tRNA splicing endonuclease assays in the high throughput screening methods of RANA, ALMSTEAD and/or RANDO, because these references specifically suggest screening small molecules libraries for compounds which disrupt tRNA interactions, including splicing, and in light of the secondary reference teaching that tRNA splicing pathway in fungi is known and analogous; and the known teaching of tRNA splicing endonuclease inhibition; with the desirability of using high throughput screening of small molecular libraries for screening enzyme binding compounds as drug candidates.